



15 FEB 2005

15 FEB 2005



INVESTOR IN PEOPLE

The Patent Office
Concept House
Cardiff Road
Newport
South Wales
NP10 8QQ

PCT/13/03/3698

REC'D 22 OCT 2003	
WIPO	PCT

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

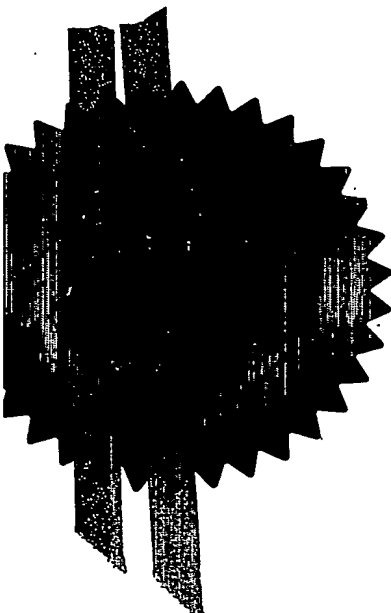
Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

PRIORITY DOCUMENT
SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH
RULE 17.1(a) OR (b)

Signed

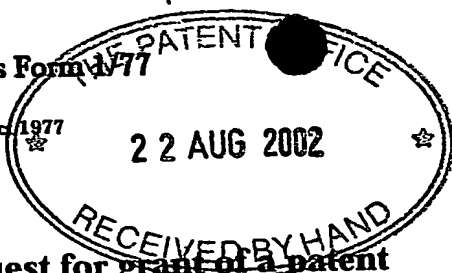
P. Mahoney

Dated 21 August 2003



An Executive Agency of the Department of Trade and Industry

BEST AVAILABLE COPY



177

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

The Patent Office

Cardiff Road
Newport
South Wales
NP10 8QQ

1. Your reference

AFB/P9192GB

23AUG02 E743172-1 D00571
01/7700 0.00-0219639.2

2. Patent application number

(The Patent Office will fill in this part)

0219639.2

22 AUG 2002

3. Full name, address and postcode of the or of each applicant (underline all surnames)

PRESTWICK SCIENTIFIC CAPITAL, INC.
1825 K Street, N.W.
Suite 1475
Washington, D.C. 20006
USA

Patents ADP number (if you know it)

8450686001

If the applicant is a corporate body, give the country/state of its incorporation

USA (Delaware)

4. Title of the invention

NOVEL PIPERIDIN-2,6-DIONE SALTS AND THEIR
USE FOR THE TREATMENT OF STRESS-RELATED
AFFECTIVE DISORDERS

5. Name of your agent (if you have one)

W. H. Beck, Greener & Co.

"Address for service" in the United Kingdom
to which all correspondence should be sent
(including the postcode)

W. H. Beck, Greener & Co.
7 Stone Buildings
Lincoln's Inn
London WC2A 3SZ

Patents ADP number (if you know it)

323001 ✓

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

Yes

- a) any applicant named in part 3 is not an inventor, or
 - b) there is an inventor who is not named as an applicant, or
 - c) any named applicant is a corporate body.
- See note (d))

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form. Do not count copies of the same document

Continuation sheets of this form

Description 10

Claim(s) 6

Abstract 1

Drawing(s)

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

Request for preliminary examination and search (*Patents Form 9/77*)

Request for substantive examination (*Patents Form 10/77*)

Any other documents
(please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature

Date

22/8/2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Mr Anthony F. Burford - (020) 7405 0921

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

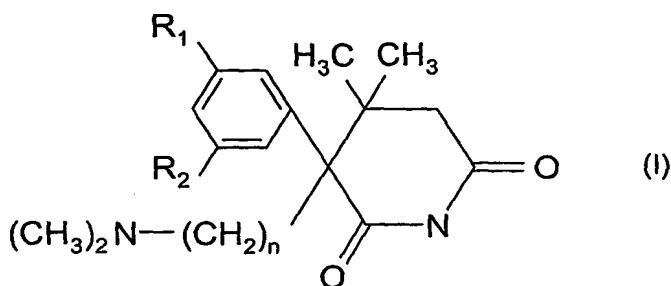
Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 08459 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

NOVEL PIPERIDIN-2,6-DIONE SALTS AND THEIR USE FOR THE
TREATMENT OF STRESS-RELATED AFFECTIVE DISORDERS.

The present invention relates to pamoate salts of certain 3-phenyl-3-
5 dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-diones and their use in the treatment of
stress-related affective disorders. The term "stress-induced affective disorder" is used
herein to include any disorder associated with elevated levels of 5-HT (5-
hydroxytryptamine; serotonin) resultant from newly synthesised 5-HT.

10 3-Phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-diones of the following
Formula I and their acid addition salts have been known since 1974 (see BE-A-808,958;
corresponding to GB-A-1,455,687 & US-A-3,963,729):



15 wherein:

R₁ represents methoxy, ethoxy or hydroxy;

R₂ represents methoxy, ethoxy, hydroxy or hydrogen; and

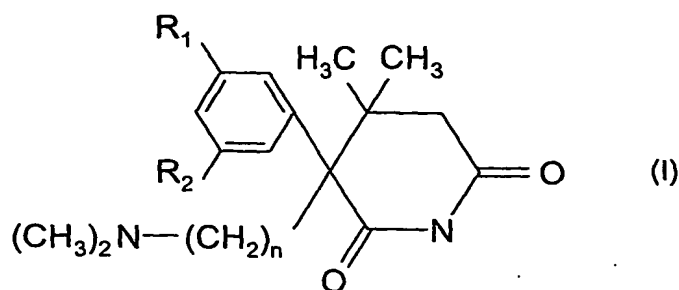
n represents 2 or 3.

20 They have been reported to have a range of pharmacological activities (see US-
A-3,963,729; US-A-4,461,771; US-A-4,738,973; US-A-4,835,151; US-A-4,835,151; US-
A-4,918,084; US-A-4,994,475; US-A-5,117,086; GB-A-2,196,251 & GB-A-2,206,491)
but were primary of interest for the treatment of stress-related affective disorders,
25 especially anxiety and depression. They are the only compounds presently known to
block selectively the activation of tryptophan hydroxylase induced by depolarisation,
metabolic inhibitors, methyl xanthine, or stress. The compound of choice for clinical
investigation was 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-
dimethylpiperidine-2,6-dione, which has been variously identified as AGN 2979 (which
30 designation will be used in this application); BTG 1501; MDL 72415 and SC 48274. A

large number of acid addition salts of AGN 2979 have been proposed but the hydrochloride has been the salt of choice because hydrochloride acid addition salts are the most commonly used acid addition salts and can be readily and inexpensively prepared and there was no reason to believe that any other salts would have any
5 advantage over the hydrochloride. There has been no previous proposal or suggestion to use a pamoate salt of AGN 2979, or of any other base of Formula I or other 3-phenyl substituted-3-dialkylaminoalkyl-4,4-dialkylpiperidin-2,6-dione, for any purpose.

A number of papers relating to clinical trials of the hydrochloride salt of AGN
10 2979 have been conducted and the results published. These showed the salt to be effective in the treatment of anxiety and depression at about 4 mg/kg/day (200-400 mg/day for human patients). However, a 1-year sub-acute toxicity study of the hydrochloride (200 mg/kg/day *p o.* (i.e. by mouth)) in rats showed that the animals suffered an immediate and continuing weight loss (40% over the 1-year period) and, as
15 revealed by post-mortem examination, hepatocyte changes which had not been detected by routine transaminase determinations during the year. As a result, the USA Food and Drugs Administration ("F.D.A") precluded the use of the dose levels previously used in the clinical trials. A subsequent clinical study by Cutler *et al* using an F.D.A. allowed dose of 1 mg b.i.d. (i.e. twice daily) (about 30 µg/kg/day) showed that the
20 hydrochloride salt of AGN 2979 possessed only marginally effective anxiolytic properties at FDA permitted dose levels.

It has now surprisingly been found that the aforementioned problems of weight loss and hepatocyte changes can be overcome by the use of the pamoate salt instead
25 of the hydrochloride, or other previously disclosed salt, of compounds of Formula I. These pamoate salts do not cause weight loss and the indications are that they will not cause hepatocyte changes over prolonged periods of treatment. Furthermore, it has been found that the pamoate salts of the compounds of Formula I, contrary to the known salts, are tasteless and allow the preparation of pharmaceutical compositions for
30 the oral administration, especially in form of suspensions, syrups and the like. Thus, according to a first aspect of the present invention, there is provided the pamoate salts of 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-diones of Formula I:



wherein:

R₁ represents methoxy, ethoxy or hydroxy;

R₂ represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

and pharmacologically acceptable solvates thereof.

Pamoic acid is 4,4'-methylenebis[3-hydroxy-2-naphthalenecarboxylic acid] and is also known as embonic acid.

The compounds of Formula I exist in optical isomers and accordingly the pamoate salts can be used in racemate form or as individual (+) or (-) isomers. Presently the (-) isomer is preferred. The salts may exist in solvated, especially, hydrated, form and may hydrate on storage in a non-airtight environment.

In a second aspect, the present invention provides methods for the treatment or prophylaxis of stress-related affective disorders which comprise administering to a human or animal patient an effective amount of a pamoate salt of a compound of Formula I or a pharmacologically acceptable solvate thereof.

In a third aspect, the present invention provides pharmaceutical compositions comprising the pamoate salt of a compound of Formula I or a pharmacologically acceptable solvate thereof and a pharmacologically acceptable diluent or carrier.

In a fourth aspect, the present invention provides the pamoate salts of compounds of Formula I and pharmacologically acceptable solvates thereof for use in treatments of the human or animal body by therapy or diagnosis practised on the human or animal body.

In a fifth aspect, the present invention provides the use of pamoate salts of compounds of Formula I and pharmacologically acceptable solvates thereof in the manufacture of medicaments for the treatment or prophylaxis of stress-related affective disorders.

5

Examples of pamoate salts of compounds of Formula I include the following:

3-(3'-methoxyphenyl)-3-(2"-N,N-dimethylaminoethyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

10 3-(3'-methoxyphenyl)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

3-(3'-hydroxyphenyl)-3-(2"-N,N-dimethylaminoethyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

3-(3'-hydroxyphenyl)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

15 3-(3'-ethoxyphenyl)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

3-(3',5'-dimethoxyphenyl)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate;

20 3-(3',5'-dihydroxy)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate; and

3-(3',5'-diethoxy)-3-(3"-N,N-dimethylaminopropyl)-4,4-dimethylpiperidin-2,6-dione pamoate.

25 The preferred pamoate salts are those of compounds of Formula I in which R₁ represents methoxy and R₂ represents methoxy or hydrogen. The most preferred salts are 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate and, especially, 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione (AGN 2979) pamoate.

30 The pamoate salts of the invention can be prepared by conventional techniques for converting a free base into an acid addition salt or for converting one acid addition salt to another. For example, the pamoate salt is prepared by treating an ethanol solution of a compound of Formula I with a cooled solution of pamoic acid in ethanol; evaporation of the solvent under reduced pressure and recrystallisation of the residue
35 from ethanol. Alternatively, a salt of a compound of Formula I may be converted into

the pamoate by neutralisation, for example with ammonium hydroxide, and subsequent treatment with pamoic acid.

The compounds of Formula I can be prepared by the processes disclosed in
5 US-A-3,963,729 or US-A-5,104,990, the disclosure of which documents are incorporated by this reference. The optical isomers can be separated in conventional manner, for example the (-) isomers can be separated by crystallisation of their (+) binaphthyl phosphoric acid salts from a suitable solvent such as ethanol.

10 The pamoate salts of the compounds of Formula I have the same qualitative pharmacological activity as that previously reported for the free base and other acid addition salts, especially the hydrochloride, and is especially useful for the treatment or prophylaxis of any stress-induced affective disorder. As mentioned above, the term "stress-induced affective disorder" is used herein to include any disorder associated with
15 elevated levels of 5-HT (5-hydroxytryptamine; serotonin) resultant from newly synthesised 5-HT. In particular, the pamoate salts can be used to treat or prevent those neurological and psychological diseases and conditions in which newly synthesised 5-HT is implicated and for which antidepressant, anxiolytic and antipsychotic drugs are presently indicated. Non-limiting examples of such diseases or conditions are
20 agoraphobia; anorexia nervosa; anxiety; anxiogenesis associated with withdrawal from drugs of abuse; bulimia nervosa; chronic paroxysmal hemicrania; depression (including prevention of depressive recurrences); diminution of the immune response associated with anxiety, depression or bereavement; disorders of sleep initiation or maintenance; disorders of the sleep-awake schedule; dream anxiety attacks; Huntington's chorea;
25 Kleine-Levin syndrome; memory disturbance; Ménière's disease, menstrual-associated sleep syndrome; migraine; motion sickness; nausea incompletely relieved by 5HT₃ antagonist administration, neurogenic pain; neuropathic pain; obsessive-compulsive disorder; panic attacks; posttraumatic stress disorder; pre-menstrual dysphoric disorder; recurrent brief depression; Restless Leg syndrome, schizophrenia; senile dementia;
30 serotonin-irritation syndrome; sleep apnoea; sleep related cardiovascular symptoms; sleep related epileptic seizures; sleep-related cluster headaches; sleep-related myoclonus syndrome; social phobia; sudden infant death syndrome; and tinnitus.

The antidepressant action of AGN 2979 pamoate is believed to result from the
35 inhibition of tryptophan hydroxylase activation, and the mechanism of this effect may

involve blockade of K⁺ channels since other metabolic inhibitors, such as guanidine and sodium cyanide, which are known to open K⁺ channels, can activate tryptophan hydroxylase and this activation can be blocked by AGN 2979 pamoate.

5 The pamoate salts of the invention can be administered in any of the manners previously proposed for the hydrochloride salt. They can be administered alone or in the form of pharmaceutical preparations to the patient being treated either orally or parenterally, for example subcutaneously or intravenously. The amount of pamoate salt administered will vary and can be any effective amount. Depending upon the patient
10 and the mode of administration, the quantity of pamoate salt administered may vary over a wide range to provide from about 0.1 mg/kg to about 20 mg/kg, usually about 0.5 mg/kg to about 10 mg/kg and preferably about 1 to about 5 mg/kg, of body weight of the patient per dose. Unit doses of these salts can contain, for example, from about 10 mg to about 500 mg, advantageously about 25 to about 200 mg. usually about 50 to about
15 100 mg of the pamoate salt and may be administered, for example, from 1 to 4 times daily. The term "unit dosage form" is used herein to mean a single or multiple dose form containing a quantity of the active ingredient in admixture with or otherwise in association with a diluent or carrier, said quantity being such that one or more predetermined units are normally required for a single therapeutic administration. In the
20 case of multiple dose forms such as liquids or scored tablets, said predetermined unit will be one fraction, such as a 5 ml (teaspoon) quantity of a liquid or a half or quarter of a scored tablet, of the multiple dose form.

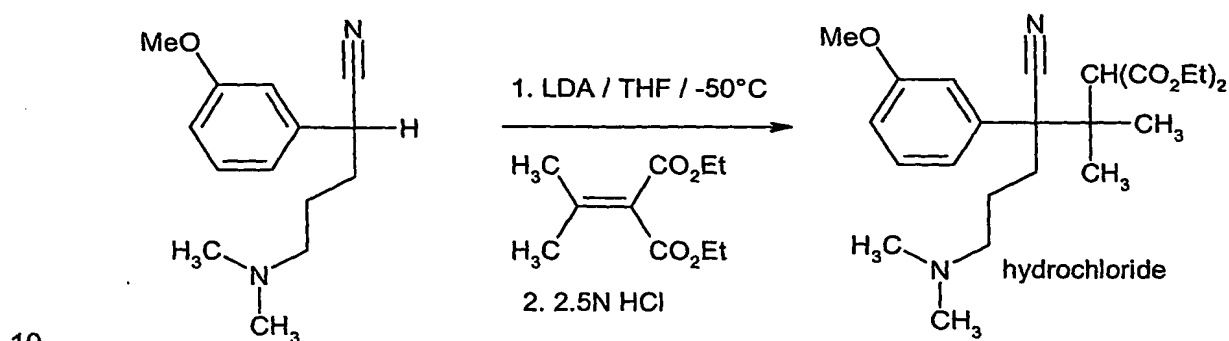
25 The pharmaceutical formulations in which form the pamoate salts of the invention will normally be utilised are prepared in a manner well known *per se* in the pharmaceutical art and usually comprise at least one active pamoate salt of the invention in admixture or otherwise in association with a pharmaceutically acceptable carrier or diluent therefor. For making those formulations, the active ingredient usually will be mixed with a carrier, or diluted by a diluent, or enclosed or encapsulated in a
30 capsule, sachet, cachet, paper or other container. A carrier or diluent may be solid, semi-solid or liquid material that serves as a vehicle, excipient or medium for the active ingredient. Suitable carriers or diluents are well known *per se*. The formulations may be adapted for enteral or parenteral use and may be administered to the patient in the form of tablets, capsules, dragees, suppositories, syrups, suspensions or the like.
35

The invention is illustrated in the following non-limiting Examples.

Example 1

Preparation of 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethyl- piperidine-2,6-dione pamoate (AGN 2979 pamoate)

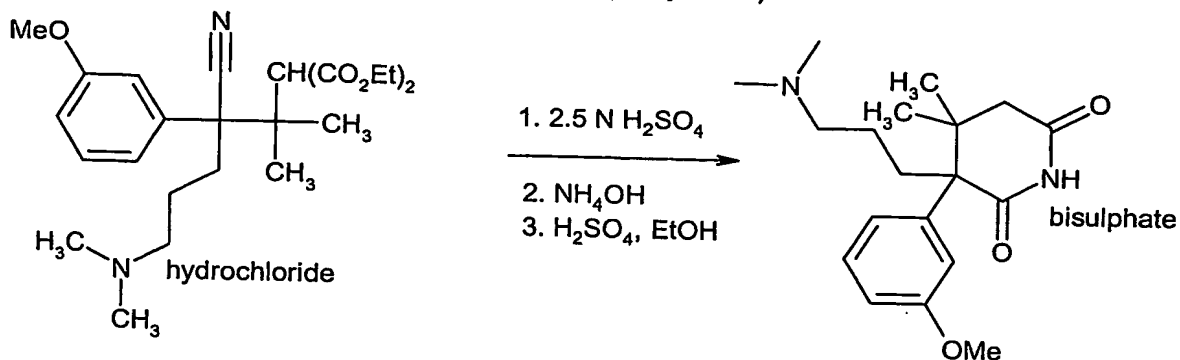
(A) Preparation of diethyl 2-[2-cyano-5-(dimethylamino)-2-(3-methoxyphenyl)-1,1-dimethylpentyl] propanedioate, monohydrochloride



A nitrogen atmosphere was applied to a reaction vessel and 50 ml of dry tetrahydrofuran is added. The solvent was cooled to less than -40°C and 32 mmoles of lithium diisopropylamide in tetrahydrofuran-heptane was added (16 ml of a 2 M solution). A solution of 6.97 g (30 mmoles) of α-[3-(dimethylamino) propyl]-3-methoxybenzeneacetonitrile in 30 ml of tetrahydrofuran was added at less than -20°C and left at this temperature for 30 min. The mixture was then cooled to -50°C and a solution of 6.62 g (33 mmoles) of diethyl isopropylidenemalonate in 30 ml of tetrahydrofuran was added to the reaction mixture at a rate such that the temperature did not exceed -50°C. The mixture was stirred at -50°C for 30 min and the cold reaction mixture added to a stirred solution of 30 ml of aqueous hydrochloric acid (36% w/w) in 100 ml of water cooled to 10°C. The mixture was extracted twice with toluene and the toluene phase is back extracted with a solution of 2 ml of hydrochloric acid (36% w/w) in 8 ml of water. The aqueous acidic extract was combined with the aqueous acidic phase from above and extracted twice with 50 ml portions of methylene chloride. The combined methylene chloride extracts were washed with water and the methylene chloride phase filtered and concentrated to low volume by distillation at atmospheric pressure. A 100 ml portion of ethyl acetate was added and the resulting slurry cooled to

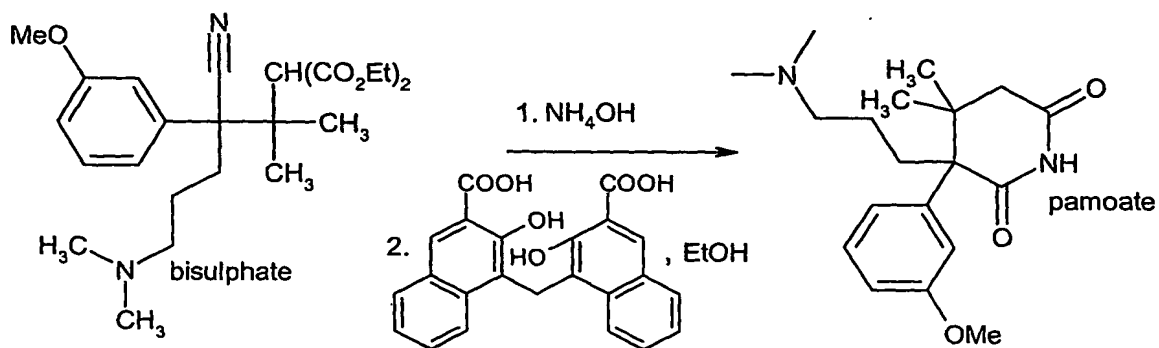
5-10°C. The resulting solid was collected by filtration, washed with ethyl acetate and dried at 50°C to give 10.1 g of white powder.

(B) Preparation of 3-(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethyl-piperidine-2,6-dione bisulphate salt (anhydrous)



A 250 ml round-bottomed flask was charged with 10 g of the above-prepared diethyl 2-[2-cyano-5-(dimethylamino)-2-(3-methoxyphenyl)-1,1-dimethylpentyl]-propanedioate mono-hydrochloride, and a solution of 10.2 g of sulphuric acid (96% w/w) in 90 ml of water was added. The reaction mixture was refluxed for about 54 hours. When the reaction was complete (as indicated by thin layer chromatography) the solution was cooled to 25°C. The aqueous solution was washed with methylene chloride, the aqueous phase mixed with methylene chloride and basified with aqueous ammonium hydroxide (29% w/w) while maintaining the temperature at less than 30°C. After separation of the layers, the aqueous phase was extracted twice with methylene chloride, the combined organic phases concentrated and the residue crystallised in tert-butyl methyl ether to give 5.7 g of white powder. The crude compound was suspended in 200 ml of absolute ethyl alcohol, 1 equivalent of concentrated sulphuric acid added and the mixture is heated under reflux for 30 minutes to dissolve the salt. After cooling, most of the solvent was evaporated under reduced pressure and the residue was by crystallised means of a 50/50 mixture of diethylether-ethyl alcohol to give 6 g of white powder (melting point = 159°-161°C) and dried under reduced pressure.

(C) Preparation of 3-(3-methoxyphenyl)-3-(3-dimethylaminopropyl]-4,4-dimethyl-piperidine-2,6-dione pamoate salt (anhydrous)



5

A solution of AGN-2979 bisulphate salt obtained in Step B (1 mmole, 430 mg) in 10 ml of water was mixed with methylene chloride (20 ml) and basified with aqueous ammonium hydroxide (29% w/w). After separation of the layers, the aqueous phase was extracted twice with methylene chloride. The combined organic phases were dried over anhydrous magnesium sulphate and the solvent was evaporated under reduced pressure. The residue was dissolved in ethanol (10 ml) and mixed with a hot solution of pamoic acid (embonic acid, 390 mg, 1 mmole) in hot ethanol (30 ml) and the mixture was heated to reflux. After cooling, the pamoate salt crystallised and the salt was recrystallised in hot ethanol to give a pale yellow powder (melting point = $146^\circ\text{-}150^\circ\text{C}$).

15

Example 2

Tablets each having the following composition are prepared by conventional tableting techniques:

20

<u>Ingredient</u>	<u>mg per tablet</u>
(a) AGN 2979 pamoate	50
(b) Lactose	51.5
(c) Maize starch dried	45
(d) magnesium stearate	1.5

Example 3

Suppositories are formed from the following composition:

<u>Ingredient</u>	<u>mg/suppository</u>
(a) AGN 2979 pamoate	20
(b) Oil of Theobroma (cocoa butter)	980

5

The compound (a) is powdered and passed through a BS No. 100 sieve (0.125 mm) and triturated with molten oil of Theobroma at 45° C. to form a smooth suspension. The mixture is well stirred and poured into moulds each of nominal 1 g capacity to produce suppositories.

10

Example 4

Pills each having the following composition are prepared by blending the active (a) and the corn starch (b), then adding the liquid glucose (c) with thorough kneading to form a plastic mass from which the pills are cut and formed:

15

<u>Ingredient</u>	<u>per pill</u>
(a) AGN 2979 pamoate	50 mg
(b) Corn starch	45 mg
(c) Liquid glucose	7 cm ³

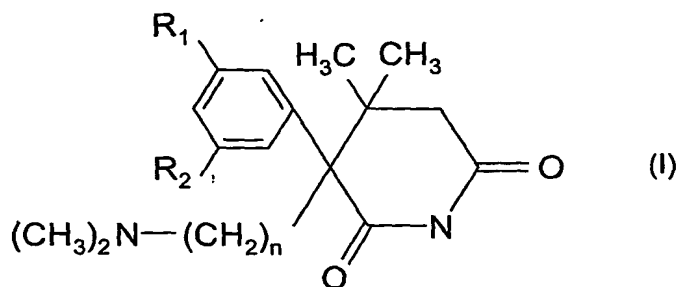
Example 5

20

Gelatine capsules each containing 50 mg AGN 2979 pamoate and 20 mg talc are prepared by passing AGN 2979 and talc separately through a fine mesh screen, mixing the sieved powders and filling the mixed powder into hard gelatine capsules at a net fill of 70 mg per capsule.

CLAIMS:

1. A pamoate salt of a 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione of Formula I:



wherein:

R₁ represents methoxy, ethoxy or hydroxy;

10 R₂ represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

or a pharmacologically acceptable solvate thereof.

2. A pamoate salt as claimed in Claim 1, wherein R₁ represents methoxy
15 and R₂ represents methoxy or hydrogen.

3. A pamoate salt as claimed in Claim 2, wherein the pamoate salt is 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

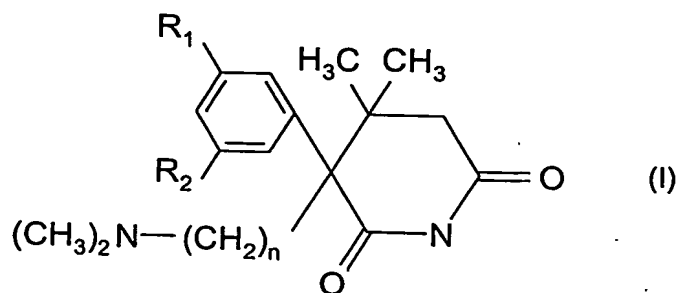
20 4. A pamoate salt as claimed in Claim 2, wherein the pamoate salt is 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

5. A pamoate salt as claimed in any one of the preceding claims, wherein
the pamoate salt is in the form of its (-) isomer.

25

6. A pamoate salt as claimed in any one of the preceding claims, wherein
the pamoate salt in the form of a hydrate.

7. A pharmaceutical composition comprising a pamoate salt of a 3-phenyl-
30 3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione of Formula I:



wherein:

R_1 represents methoxy, ethoxy or hydroxy;

R_2 represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

or a pharmacologically acceptable solvate thereof

and a pharmacologically acceptable diluent or carrier.

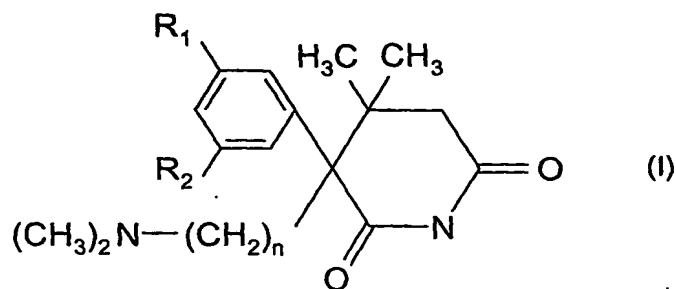
8 A pharmaceutical composition as claimed in Claim 7 for the treatment or prophylaxis of a stress-related affective disorder.

9 A pharmaceutical composition as claimed in Claim 7 or Claim 8, wherein the 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione is as defined in any one of Claims 2, 5 and 6.

10. A pharmaceutical composition as claimed in Claim 9, wherein the pamoate salt is 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

11. A pharmaceutical composition as claimed in Claim 9, wherein the pamoate salt is 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

12. A pamoate salt of a 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione of Formula I:



wherein:

R_1 represents methoxy, ethoxy or hydroxy;

R_2 represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

or a pharmacologically acceptable solvate thereof

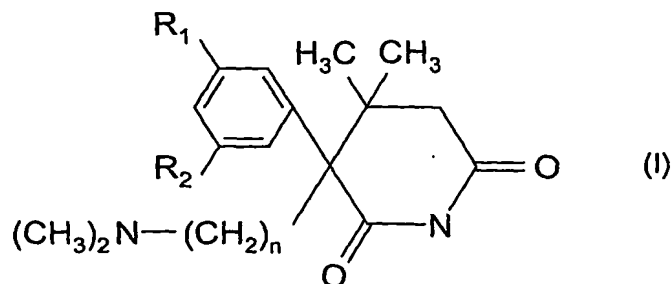
for use in treatment of the human or animal body by therapy or diagnosis practised on the human or animal body.

13 A pamoate salt as claimed in Claim 12, wherein the 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione is as defined in any one of Claims 2, 5 and 6.

14. A pamoate salt as claimed in Claim 13, wherein the pamoate salt is 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

15. A pamoate salt as claimed in Claim 13, wherein the pamoate salt is 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

16. Use of a pamoate salt of a 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione of Formula I:



wherein:

R₁ represents methoxy, ethoxy or hydroxy;

R₂ represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

- 5 or a pharmacologically acceptable solvate thereof
in the manufacture of a medicament for the treatment or prophylaxis of stress-related
affective disorders.

10 17. The use as claimed in Claim 16, wherein the stress-related affective
disorder is selected from agoraphobia; anorexia nervosa; anxiety; anxiogenesis
associated with withdrawal from drugs of abuse; bulimia nervosa; chronic paroxysmal
hemicrania; depression (including prevention of depressive recurrences); diminution of
the immune response associated with anxiety, depression or bereavement; disorders of
sleep initiation or maintenance; disorders of the sleep-awake schedule; dream anxiety
15 attacks; Huntington's chorea; Kleine-Levin syndrome; memory disturbance; Ménière's
disease, menstrual-associated sleep syndrome; migraine; motion sickness; nausea
incompletely relieved by 5HT₃ antagonist administration, neurogenic pain; neuropathic
pain; obsessive-compulsive disorder; panic attacks; posttraumatic stress disorder; pre-
menstrual dysphoric disorder; recurrent brief depression; Restless Leg syndrome,
20 schizophrenia; senile dementia; serotonin-irritation syndrome; sleep apnoea; sleep
related cardiovascular symptoms; sleep related epileptic seizures; sleep-related cluster
headaches; sleep-related myoclonus syndrome; social phobia; sudden infant death
syndrome; and tinnitus.

25 18. The use as claimed in Claim 17, wherein the medicament is for the
treatment or prophylaxis of anxiety.

19. The use as claimed in Claim 17, wherein the medicament is for the
treatment or prophylaxis of depression.

30 20. The use as claimed in Claim 17, wherein the medicament is for the
treatment or prophylaxis of migraine.

21. The use as claimed in Claim 17, wherein the medicament is for the
35 treatment or prophylaxis of sleep apnoea.

22. The use as claimed in any one of Claims 16 to 21, wherein the 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione is as defined in any one of Claims 2, 5 and 6.

5

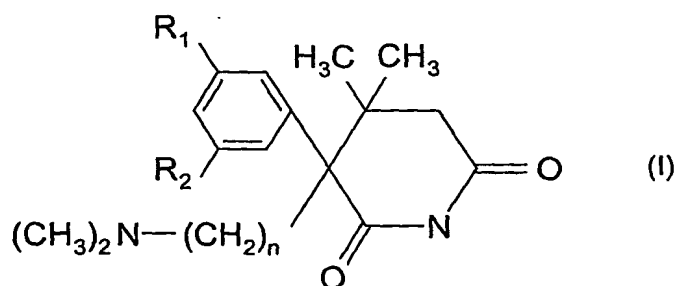
23. The use as claimed in Claim 22, wherein the pamoate salt is 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

24. The use as claimed in Claim 22, wherein the pamoate salt is 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

10

25. A method for the treatment or prophylaxis of a stress-related affective disorder which comprises administering to a human or animal patient an effective amount of a pamoate salt of a 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione of Formula I:

15



wherein:

20

R_1 represents methoxy, ethoxy or hydroxy;

R_2 represents methoxy, ethoxy, hydroxy or hydrogen; and

n represents 2 or 3,

or a pharmacologically acceptable solvate thereof.

26. A method as claimed in Claim 25, wherein the stress-related affective disorder is selected from the group consisting of agoraphobia; anorexia nervosa; anxiety; anxiogenesis associated with withdrawal from drugs of abuse; bulimia nervosa; chronic paroxysmal hemicrania; depression (including prevention of depressive recurrences); diminution of the immune response associated with anxiety, depression or bereavement; disorders of sleep initiation or maintenance; disorders of the sleep-awake

30

schedule; dream anxiety attacks; Huntington's chorea; Kleine-Levin syndrome; memory disturbance; Ménière's disease, menstrual-associated sleep syndrome; migraine; motion sickness; nausea incompletely relieved by 5HT₃ antagonist administration, neurogenic pain; neuropathic pain; obsessive-compulsive disorder; panic attacks; 5 posttraumatic stress disorder; pre-menstrual dysphoric disorder; recurrent brief depression; Restless Leg syndrome, schizophrenia; senile dementia; serotonin-irritation syndrome; sleep apnoea; sleep related cardiovascular symptoms; sleep related epileptic seizures; sleep-related cluster headaches; sleep-related myoclonus syndrome; social phobia; sudden infant death syndrome; and tinnitus.

10

27. A method as claimed in Claim 26, wherein the stress-related affective disorder is anxiety.

28. A method as claimed in Claim 26, wherein the stress-related affective 15 disorder is depression.

29. A method as claimed in Claim 26, wherein the stress-related affective disorder is migraine.

20 30. A method as claimed in Claim 26, wherein the stress-related affective disorder is sleep apnoea.

31. A method as claimed in any one of Claims 25 to 30, wherein the 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-dione is as defined in any one of 25 Claims 2, 5 and 6.

32. A method as claimed in Claim 31, wherein the pamoate salt is 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

30 33. A method as claimed in Claim 31, wherein the pamoate salt is 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

ABSTRACT

NOVEL PIPERIDIN-2,6-DIONE SALTS AND THEIR USE FOR THE
TREATMENT OF STRESS-RELATED AFFECTIVE DISORDERS.

5

Novel pamoate salts of certain 3-phenyl-3-dimethylaminoalkyl-4,4-dimethylpiperidin-2,6-diones and pharmacologically acceptable solvates thereof are devoid of the weight loss and hepatocyte changes in the rat which limited to marginally effective levels the permitted clinical doses of the corresponding hydrochlorides in the treatment or prophylaxis of stress-related affective disorders such as anxiety, depression, migraine and sleep apnoea. The preferred pamoate salts are 3(3,5-dimethoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate and, especially, 3(3-methoxyphenyl)-3-(3-dimethylaminopropyl)-4,4-dimethylpiperidine-2,6-dione pamoate.

10

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☒ FADED TEXT OR DRAWING
- ☒ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.